MEIOTIC DISORDERS INDUCED BY NADOLOL IN ALLIUM CEPA L.

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Several investigators studied the effect of drugs on either somatic or germ cells among them, Shahab et al. (1983, 1985), Shehab and Abo El-Kheir (1984), Reddy and Subramanyam (1985) and El-Bayoumi et al. (1985).

The present investigation was undertaken to elucidate the effects of Nadolol drug on meiosis of Allium cepa L. pollen mother cells (P.M.C.s).

Nadolol is used in the treatment of hypertension and angina pectoris. In hypertension the initial starting dose is usually 40 to 80 μg once daily. The usual maintainance dose is 80 to 320 μg once daily.

MATERIAL AND METHODS

Allium cepa (Var. Giza 5) flower buds were treated for 3 and 6 hours with the different concentrations of the drug (50, 100 µg/ml and 1 mg/ml) using a piece of cotton soaked with the drug solution. The treated flower buds were collected 24 and 48 hours after treatment (recovery test) at random from 20 plants for each treatment. Tap water was used for control in the same manner. Flower buds were fixed in Carnoy's fluid and examined using aceto carmine smear method.

RESULTS AND DISCUSSION

Tables 1 and 2 show that the drug induced a high percentage of aberrations in all treatments of the drug. This percentage increased with the increase of concentration in all treatments of 24 & 48 hours recovery, except 3 hrs treatment with 1 mg/ml after 24 hrs. recovery.

The effect of Nadolol drug was permanent, since the percentage of aberration increased with lapse of time of recovery. After 24 & 48 hrs. recovery the first division has the highest percentage of anomalis except 6 hrs. treatment with 100 µg after 24 hrs. recovery (Table 1) and 3 & 6 hrs. treatment with 50 µg/ml after 48 hrs. recovery (Table 2). The decrease in aberration percentage with the progress of meiotic stages from the first to the second division may indicate the loss (elimination) of aberrant cells (gametes) from the population, and not the recovery of these cells from aberration events Ashour (1988).

No trend was observed between the percentage of anomalis and time of treatment in the second division.

Tables 3 and 4 represent the percentage of the different types of abnormalities induced as a result of treatment of \underline{Allium} \underline{cepa} flower buds with the different concentrations of Nadolol drug.

Stickiness was the most prominent abnormality in all treatments of the drug. The highest percentage of stickiness was observed in diakinesis and M1 and it gradually decreased in the later meiotic division. Fig.1 shows sticky MI with swallen and grouping of bivalents. The sticky clumped chromosomes when try to separate they form sticky bridges (Fig. 2).

A less dominant abnormality was the disturbed. Fig. 3 shows disturbed anaphase 1 and Fig. 4 shows the disturbance in one pole only of anaphase 1.

Nadolol induced clastogenic effect such as laggards (Fig. 5) and fragment and bridges Fig. 6.

In addition to above mentioned abnormalities despiralization assynchronization, diagonal and univalent were observed in low percentage in some of the treatments.

Micro and multinucleate cells were observed which are a result of either spindle disturbance or lagging chromosomes. The same types of abnormality have been met with and discussed by a number of authors, among them Kunzel and Mirslaw 1966, Vig 1969 and Bezo et al., 1980.

SUMMARY

In this investigation the effect of nadolol on $\underbrace{Allium}_{\text{cepa}}$ PMCs was studied. The drug gave high percentage of abnormalities. The percentage of abnormalities was higher in the 1st division. Metaphases have the highest percentage of aberrations.

Different types of abnormalities were met with stickiness, disturbed metaphases and ana-telophases, lagging chromosomes, bridges, fragments and despiralization. Micro and multinucleate cells were also recorded.

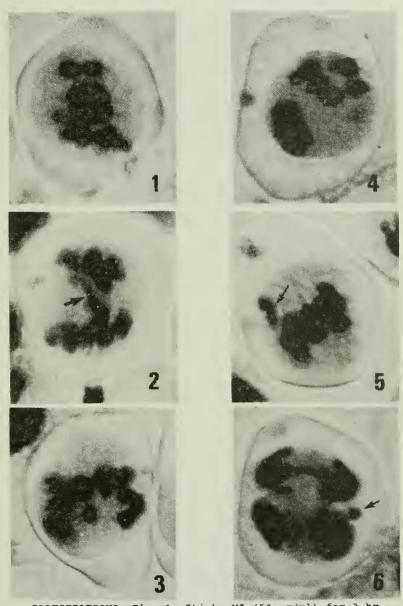
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ILLUSTRATIONS: Fig. 1: Sticky MI (50 μg/ml) for 3 hr nadolol after 24 hr recovery. Fig. 2: AI with bridge (50 μg/ml) for 6 hr nadolol after 24 hr recovery. Fig. 3: Disturbed AI (1 mg/ml) for 6 hr nadolol after 48 hr recovery. Fig. 4: Disturbed AI at one pole only (100 μg/ml) for 3 hr nadolol after 24 hr recovery. Fig. 5: Lagging chromosome at MI (1 mg/ml) for 3 hr nadolol after 48 hr recovery. Fig. 6: AII with bridge and fragment (50 μg/ml) for 3 hr nadolol after 48 hr M = Metaphase; A = Anaphase

Percentage of total abnormalities and abnormalities in each phase of meiosis in Allium cepal. treated with the drug after 24 hours recovery. Table (1):

		No. of	Jo oN	Jo %		lst di	division		2nd c	2nd division		
Ð	Experimental agent	Count- ed cells		total abn.	% of abn. metaphase	% of abn . anapha <i>s</i> e.	% of abn. telophase	Total % of abn.	% of abn. metaphase	% of % of abn. metaphase	% of abn. te lophase	Total % of abn.
	Control	4787	ħ	2.38	5.36	2.27	0.7	4.28	0.43	9.1	0.63	1.12
	50 µg/ml.	3581	709	19.80	29.4	53.54	20.00	25.87	19.17	21.26	8.15	15.46
ponts	100 µg/ml.	2356	729	30.94	35.16	6.09	37.92	37.52	75.76	34.85	10.07	22.90
٤	1 mg/ml.	4182	948	20.23	27.34	75.00	16.59	23.99	46.25	52.14	5.75	18.17
	50 µg/ml.	3431	630	18.36	36.08	37.70	82.6	22.96	27.67	39.46	5.43	14.76
sanoi	100 pg/ml.	2464	532	21.59	20.71	29.48	11.71	19.04	29.36	29.75	11.52	23.59
1 7	I mg/mil.	3086	797	24.85	33.68	49.49	12.04	51.9	51.9	41.15	1.32	15.42
;												

No = Number Abn = Abnormal

20 | | | | | | | | | | | |

		No. of	No. of	50 %		I st division	ion			2nd division	vision	
Ex	Experimental agent	counted	abn.	total abn.	% of abn. metaphase	% of abn. anaphase	% of abn. telophase	Total % of abn.	% of abn.	% of abn.	% of abn. telophase	Total % of abn.
	Control	4375	153	3.50	7.15	2.19	1.10	3.81	3.90	3.45	2.03	3.04
	50 µg/ml.	1549	358	23.11	34.15	30.56	9.14	26.54	32.27	36.80	4.25	54.19
sanou	100 µg/ml.	1872	621	33.17	45.83	35.71	36.81	38.78	38.84	34.71	14.66	28.50
1 8	1 mg/ml.	2011	725	36.05	47.49	39.76	33.82	41.57	44.29	40.87	21.44	31.30
	50 µg/ml.	8061	788	41.30	41.82	48.65	35.02	39.38	50.58	47.81	25.27	42.52
sano	100 µg/ml.	6111	483	43.16	51.72	51.02	45.46	48.89	53.82	59.13	16.85	39.31
19	I mg/ml.	1387	620	44.70	59.54	57.14	42.98	55.77	51.42	10.64	21.16	37.14

Percentage of each type of abnormalities in Allium cepa L.P.M.Cs. treated with 50,100 µg/ml and I mg/ml Nadolol after 24 hours recovery. Table (3):

Stickiness Disturbins lagging bridge fragment diagonal univalent micronuclei multinucleate assynchro- despira-	- 0.88 1.75	0.56 13.68 5.08 0.42 0.71	- 20.85 0.69 0.27 -	0.95 9.10 8.51 0.35 -	- 8.89 4.26 0.48 -	- 15.04 1.13 0.19 0.19	- 3.13 0.65 0.13 1.30
nt diagonal un	1	1.4.1	3.57	0.70	1.43	4.70	1.57
bridge fragme	0.88	9.03 11.14 5.78	8.92 10.70	3.31 3.43	7.46 1.75	4.51 9.02	6.00 15.25 4.69
lagging	10.53	9.03	6.31	8.87	8.41	7.14	00.9
Disturbins	7.01	12.83	3.98	14.54	13.49	11.28	12.65
Stickiness	78.95	39.35	44.72	50.24	53.81	8.94	54.63
Experimental agents	Control	50 µg/ml. 39.35	100 µg/ml. 44.72	I mg/ml. 50.24	50 µg/ml.	100 µg/ml. 46.8	l mg/ml. 54.63

Percentage of each type of abnormalities of Allium cepa L.P.M.Cs. treated with 50, 100 µg/ml, 1 mg/ml Nadolol after 48 hours recovery during the 3 and 6 hours exposure. Table (4):

											asswuchro-	despira-
age	Experimental agent	stickiness	disturbins	lagging	bridge	fragment	diagonal	univalent	micronuclei	stickiness disturbins lagging bridge fragment diagonal univalent micronuclei multinucleate nization lisation	nization	lisation
ŭ	Control	71.90	94.01	9.15	2.61	ı	ı	ı	1.96	3.92	1	1
	50 µg/ml.	49.64	12.29	5.87	10.89	10.62	2.51	1.68	2.79	3.35	0.56	
ponts	100 µg/ml. 42.51	42.51	10.31	8.21	13.21	4.35	0.16	1.77	18.68	0.64	0.16	,
_	I mg/ml.	37.93	9.38	11.17	11.17 17.66	4.41	1.38	1	14.90	2.35	0.14	89.0
	50 µg/ml.	37.18	12.06	9.52	9.52 10.15	6.22	1.78	0.51	10.03	12.18	0.25	0.13
yonus	100 µg/ml.	38.92	7.87	4.97	5.38	11.39	2.28	1.45	19.61	7.45	0.62	1
9	I mg/ml.	50.32	9.52	5.97	5.00	6.29	0.97	1.13	0.48	20.00	ı	0.32
						The Market						